L1

L2 L3

L4

=>

(FILE 'HOME' ENTERED AT 09:00:56 ON 02 SEP 2005)

FILE 'DISSABS, 1MOBILITY, AGRICOLA, AQUASCI, BIOTECHNO, COMPENDEX, COMPUAB, CONF, CONFSCI, ELCOM, HEALSAFE, IMSDRUGCONF, LIFESCI, OCEAN, PAPERCHEM2, PASCAL, POLLUAB, SOLIDSTATE, ADISCTI, ADISINSIGHT, ADISNEWS, ANABSTR, ANTE, AQUALINE, BIOBUSINESS, BIOCOMMERCE, 'ENTERED AT 09:01:07 ON 02 SEP 2005

1690 S NOPE

122 S L1 AND (VENTRICULAR ZONE OR DCC OR PUNC OR NCAM OR AXONAL OR

68 S L2 AND (FUNCTION OR ACTIVITY)

67 DUP REM L3 (1 DUPLICATE REMOVED)

ANSWER 19 OF 67 DGENE COPYRIGHT 2005 The Thomson Corp on STN DGENE ACCESSION NUMBER: AAU77405 Protein Novel cytoplasmic, nuclear, membrane bound and secreted NOVX TITLE: polypeptides, useful for treating developmental disorders, endocrine disorders, vascular disorders, infectious diseases and neurodegenerative disorders -Rastelli L; Shimkets R A; Zerhusen B; Malyankar U M; Padigaru **INVENTOR:** (CURA-N) CURAGEN CORP. PATENT ASSIGNEE: WO 2002006329 A2 20020124 178 PATENT INFO: APPLICATION INFO: WO 2001-US22709 20010718 US 2000-218870P PRIORITY INFO: 20000718 US 2000-218875P 20000718 US 2000-218901P 20000718 US 2000-220273P 20000724 US 2000-220912P 20000726 US 2000-221233P 20000727 US 2000-221650P 20000728 DOCUMENT TYPE: Patent LANGUAGE: English 2002-179781 [23] OTHER SOURCE: CROSS REFERENCES: N-PSDB: ABK11101 Human NOV1 protein, homologue of NOPE/PUNC DESCRIPTION: Iq proteins. The present invention relates to the isolation of novel human AΒ polypeptides referred to as NOV1, NOV2, NOV3, NOV4a, NOV4b, NOV5A, NOV5b, NOV6 AND NOV7, and the polynucleotide sequences encoding them. The NOVX polypeptides are related to NOPE, cadherin, interferon alpha-13, ADAM, ankyrin repeat-containing, transpanin or semaphorin polypeptides. The sequences of the invention are useful for identifying

The present invention relates to the isolation of novel human polypeptides referred to as NOV1, NOV2, NOV3, NOV4a, NOV4b, NOV5A, NOV5b NOV6 AND NOV7, and the polynucleotide sequences encoding them. The NOVX polypeptides are related to NOPE, cadherin, interferon alpha-13, ADAM, ankyrin repeat-containing, transpanin or semaphorin polypeptides. The sequences of the invention are useful for identifying an agent (a cellular receptor or downstream effector) that binds to the NOVX polypeptide, or an agent that modulates it's expression or activity. They are useful for treating or preventing NOVX-associated disorders such as developmental disorders, endocrine disorders, vascular diseases, gastrointestinal disorders, respiratory disorders, inflammatory disorders, blood disorders, reproductive disorders, neurodegenerative disorders, autoimmune and immune disorders, infectious diseases, cardiovascular disorders, cancers, and other disorders related to cell signal processing and metabolic pathway modulation. The present sequence represents the human NOV1 protein.

DESC Human NOV1 protein, homologue of NOPE/PUNC Ig

L4 ANSWER 31 OF 67 DGENE COPYRIGHT 2005 The Thomson Corp on STN

ACCESSION NUMBER: AAE05252 Protein DGENE

TITLE: Murine Nope polypeptides and nucleic acids useful

for preventing, diagnosing and treating colonic cancer and

Bardet-Biedl syndrome -

INVENTOR: Salbaum J M

PATENT ASSIGNEE: (NEUR-N) NEUROSCIENCES RES FOUND INC.

PATENT INFO: WO 2001049714 A2 20010712 99

APPLICATION INFO: WO 2000-US29698 20001026 PRIORITY INFO: US 2000-174496 20000104 US 2000-205789 20000519

DOCUMENT TYPE: Patent
LANGUAGE: English
OTHER SOURCE: 2001-44

OTHER SOURCE: 2001-441846 [47] CROSS REFERENCES: N-PSDB: AAD10022

extracellular domain.

DESCRIPTION: Mouse Nope (neighbour of punc ell)

extracellular domain.

AΒ The present invention relates to Nope (neighbour of punc ell) which is used in the prevention, treatment and diagnosis of diseases associated with inappropriate Nope expression such as cancers especially colonic cancer and genetic disorders, as Nope is thought to be a tumour suppressor. Nope gene is located on chromosome 9 and is used in gene therapy. Nope is used as vaccine. Nope gene may be administered to treat diseases by rectifying mutations or deletions in a patient's genome that affect the activity of Nope by expressing inactive proteins or to supplement the patients own production of Nope polypeptides. Nope gene is used to study the expression and function of Nope polypeptides and their role in metabolism through the creation of transgenic animal models. The anti-Nope antibodies and Nope antagonists may also be used to down regulate Nope expression and activity for the treatment of Bardet-Biedl syndrome which is an autosomal recessive disorder characterised by mental retardation, obesity, polydactyly, retinitis pigmentosa and hypogonadism. Patients with Bardet-Biedl syndrome have a high incidence of hypertension, diabetes mellitus and renal and cardiovascular anomalies. The present sequence is mouse Nope (neighbour of punc ell)

TI Murine Nope polypeptides and nucleic acids useful for preventing, diagnosing and treating colonic cancer and Bardet-Biedl syndrome -

DESC Mouse Nope (neighbour of punc ell) extracellular domain.

Mouse; Nope; neighbour of punc ell; cytostatic; neuroprotective; vaccine; gene therapy; cerebroprotective; colonic cancer; mental retardation; tumour suppressor; chromosome 9; transgenic animal; genetic disorder; obesity;. . .

AΒ The present invention relates to Nope (neighbour of punc ell) which is used in the prevention, treatment and diagnosis of diseases associated with inappropriate Nope expression such as cancers especially colonic cancer and genetic disorders, as Nope is thought to be a tumour suppressor. Nope gene is located on chromosome 9 and is used in gene therapy. Nope is used as vaccine. Nope gene may be administered to treat diseases by rectifying mutations or deletions in a patient's genome that affect the activity of Nope by expressing inactive proteins or to supplement the patients own production of Nope polypeptides. Nope gene is used to study the expression and function of Nope polypeptides and their role in metabolism through the creation of transgenic animal models. The anti-Nope antibodies and Nope antagonists may also be used to down regulate Nope expression and activity for the treatment of Bardet-Biedl syndrome which is an autosomal recessive disorder characterised by mental retardation, obesity, polydactyly, retinitis pigmentosa. . . Bardet-Biedl syndrome have a high incidence of hypertension, diabetes mellitus and renal and cardiovascular anomalies. The present sequence is mouse Nope

```
ANSWER 32 OF 67 DGENE COPYRIGHT 2005 The Thomson Corp on STN
ACCESSION NUMBER: AAE05251 Protein
                                          DGENE
                  Murine Nope polypeptides and nucleic acids useful
TITLE:
                  for preventing, diagnosing and treating colonic cancer and
                  Bardet-Biedl syndrome -
INVENTOR:
                  Salbaum J M
PATENT ASSIGNEE: (NEUR-N) NEUROSCIENCES RES FOUND INC.
                                                             99
PATENT INFO:
                 WO 2001049714 A2 20010712
APPLICATION INFO: WO 2000-US29698 20001026
PRIORITY INFO: US 2000-174496
                                      20000104
                US 2000-205789
                                      20000519
DOCUMENT TYPE: Patent
LANGUAGE:
                English
OTHER SOURCE:
                 2001-441846 [47]
CROSS REFERENCES: N-PSDB: AAD10021
                 Mouse Nope (neighbour of punc ell)
DESCRIPTION:
                  protein.
AB
      The present invention relates to Nope (neighbour of
      punc ell) which is used in the prevention, treatment and
      diagnosis of diseases associated with inappropriate Nope
      expression such as cancers especially colonic cancer and genetic
      disorders, as Nope is thought to be a tumour suppressor.
      Nope gene is located on chromosome 9 and is used in gene therapy.
      Nope is used as vaccine. Nope gene may be administered
      to treat diseases by rectifying mutations or deletions in a patient's
      genome that affect the activity of Nope by expressing
      inactive proteins or to supplement the patients own production of
      Nope polypeptides. Nope gene is used to study the
      expression and function of Nope polypeptides and
      their role in metabolism through the creation of transgenic animal
      models. The anti-Nope antibodies and Nope antagonists
      may also be used to down regulate Nope expression and
      activity for the treatment of Bardet-Biedl syndrome which is an
      autosomal recessive disorder characterised by mental retardation,
      obesity, polydactyly, retinitis pigmentosa and hypogonadism. Patients
      with Bardet-Biedl syndrome have a high incidence of hypertension,
      diabetes mellitus and renal and cardiovascular anomalies. The present
      sequence is mouse Nope (neighbour of punc ell)
      protein.
TI
      Murine Nope polypeptides and nucleic acids useful for
      preventing, diagnosing and treating colonic cancer and Bardet-Biedl
      syndrome -
DESC
      Mouse Nope (neighbour of punc ell) protein.
KW
      Mouse; Nope; neighbour of punc ell; cytostatic;
      neuroprotective; vaccine; gene therapy; cerebroprotective; colonic
      cancer; mental retardation; tumour suppressor; chromosome 9; transgenic
      animal; genetic disorder; obesity;.
AΒ
      The present invention relates to Nope (neighbour of
      punc ell) which is used in the prevention, treatment and
      diagnosis of diseases associated with inappropriate Nope
      expression such as cancers especially colonic cancer and genetic
      disorders, as Nope is thought to be a tumour suppressor.
      Nope gene is located on chromosome 9 and is used in gene therapy.
      Nope is used as vaccine. Nope gene may be administered
      to treat diseases by rectifying mutations or deletions in a patient's
      genome that affect the activity of Nope by expressing
      inactive proteins or to supplement the patients own production of
      Nope polypeptides. Nope gene is used to study the
      expression and function of Nope polypeptides and
      their role in metabolism through the creation of transgenic animal
      models. The anti-Nope antibodies and Nope antagonists
      may also be used to down regulate Nope expression and
      activity for the treatment of Bardet-Biedl syndrome which is an
      autosomal recessive disorder characterised by mental retardation,
      obesity, polydactyly, retinitis pigmentosa. . . Bardet-Biedl syndrome
     have a high incidence of hypertension, diabetes mellitus and renal and
      cardiovascular anomalies. The present sequence is mouse Nope
```